

Supporting Information

Metallic Ion Leaching from Heterogeneous Catalysts: An Overlooked Effect in the Study of Catalytic Ozonation Processes

Submitted by

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17 This PDF file includes:

18 Table S1

19 Figure S1 to S7

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26 Table S1. HPLC conditions for OA, nitrobenzene, *p*CBA measurements

Compound	Wavelength	Mobile phase	Flow rate	Injection volume
	(nm)		(mL/min)	(μ L)
Oxalic acid	268	20 mM KH ₂ PO ₄ /CH ₃ OH (97:3)	1	10
Nitrobenzene	254	DI/CH ₃ OH (50:50)	1	10
<i>p</i> CBA	234	20 mM H ₃ PO ₄ /CH ₃ OH (30:70)	1	10

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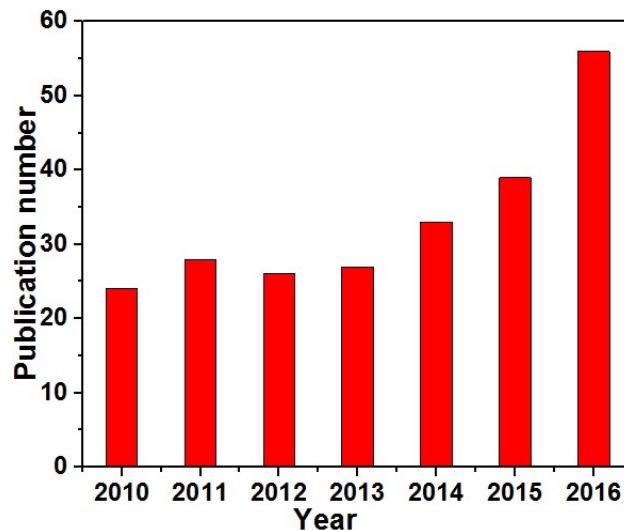
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45 Figure S1. Number of research articles in the field of heterogeneous catalytic ozonation
46 published from 2010 to 2016 according to Scopus (Elsevier).

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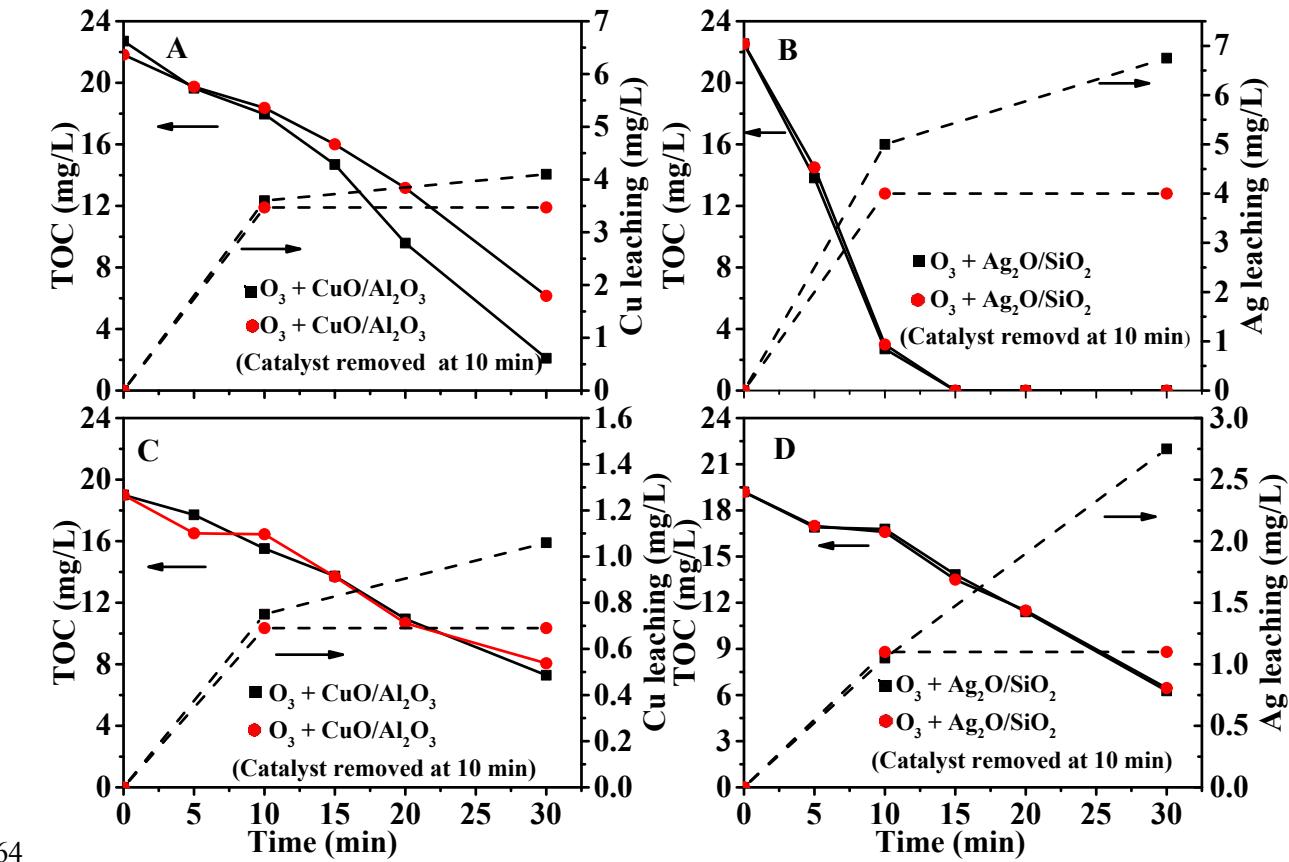
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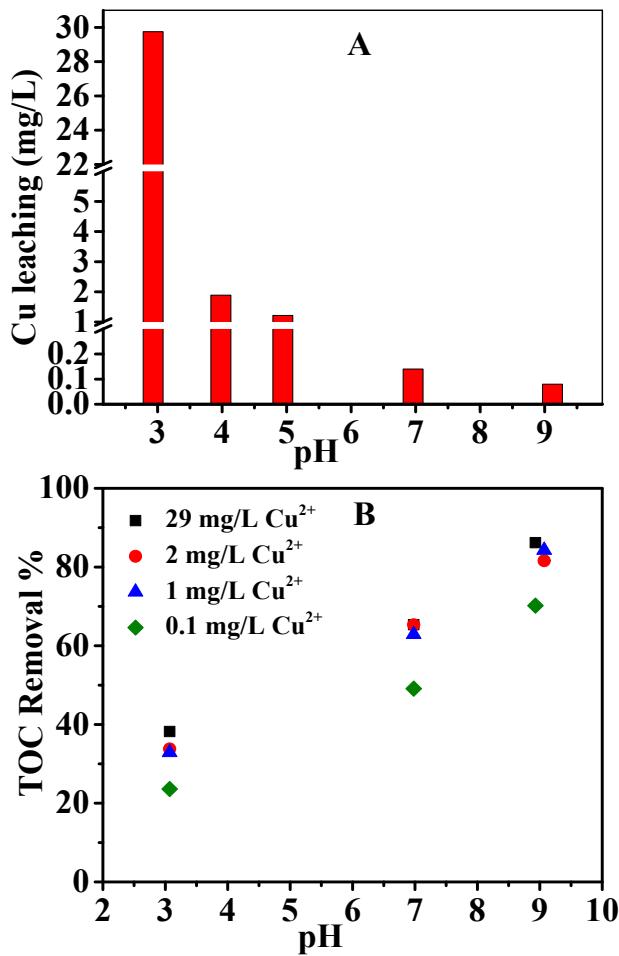
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65 Figure S2. Mineralization of models compounds during catalytic ozonation: OA (A & B); and
66 Nitrobenzene (C & D). [Initial OA: 1 mM; initial nitrobenzene: 0.325 mM; working solution:
67 200 mL; catalyst dose: 1 g].

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70 Figure S3. (A) Cu leaching in buffered nitrobenzene solutions at different pH [catalyst dose: 1 g
 71 CuO/SiO₂; nitrobenzene concentration: 0.325 mM; working solution: 200 mL; stirring time: 30
 72 min]; (B) TOC reduction at different pH and copper ion concentrations. [Initial nitrobenzene
 73 concentration: 0.325 mM; working solution: 200 mL; reaction time: 30 min].

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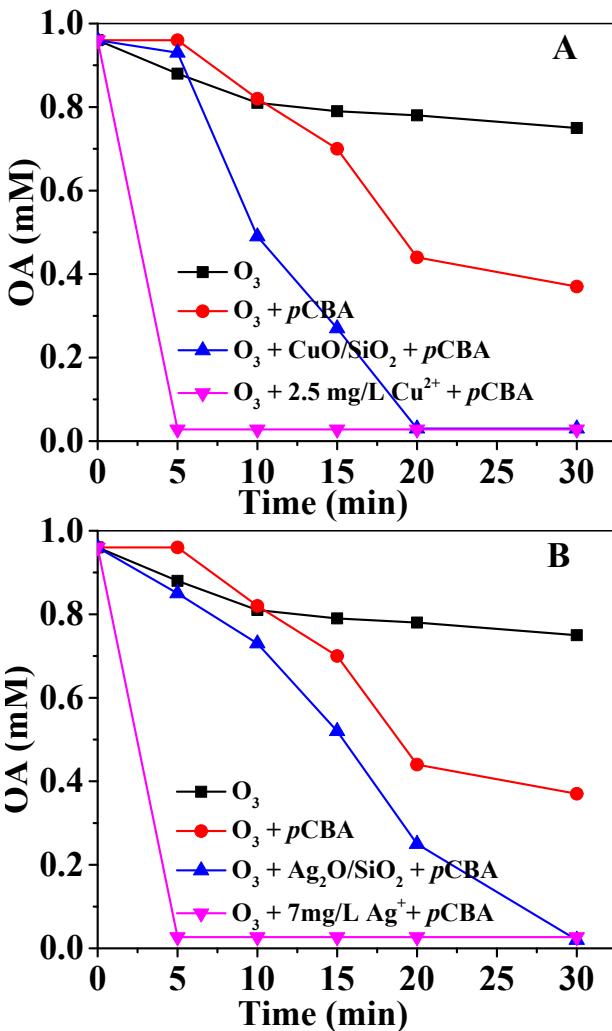
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84 Figure S4. Effect of *p*CBA on the degradation of OA. [Initial OA: 1.0 mM; working solution:
85 200 mL; catalyst dose: 1 g; *p*CBA: 18 mg/L].

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91 Figure S5. Comparison of TOC reduction with and without *p*CBA addition. [Initial OA: 1.0 mM;
92 working solution: 200 mL; catalyst dose: 1 g; reaction time: 30 min; initial *p*CBA concentration:
93 18 mg/L].

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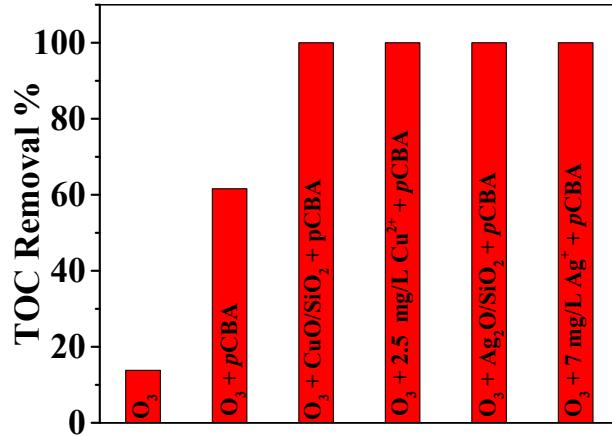
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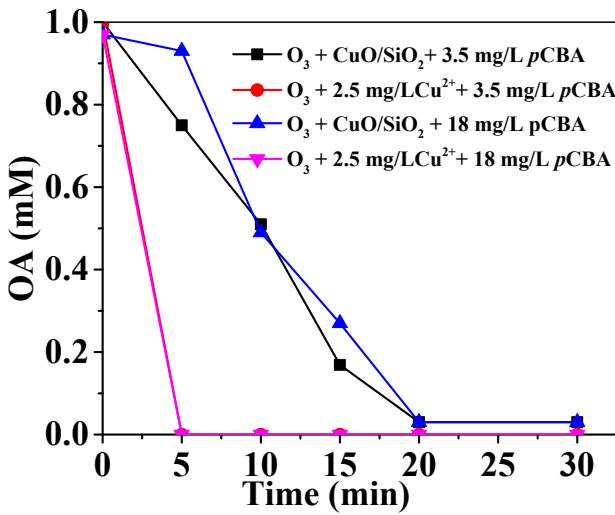
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113 Figure S6. Effect of *p*CBA concentrations on the degradation of OA. [Initial OA: 1.0 mM;
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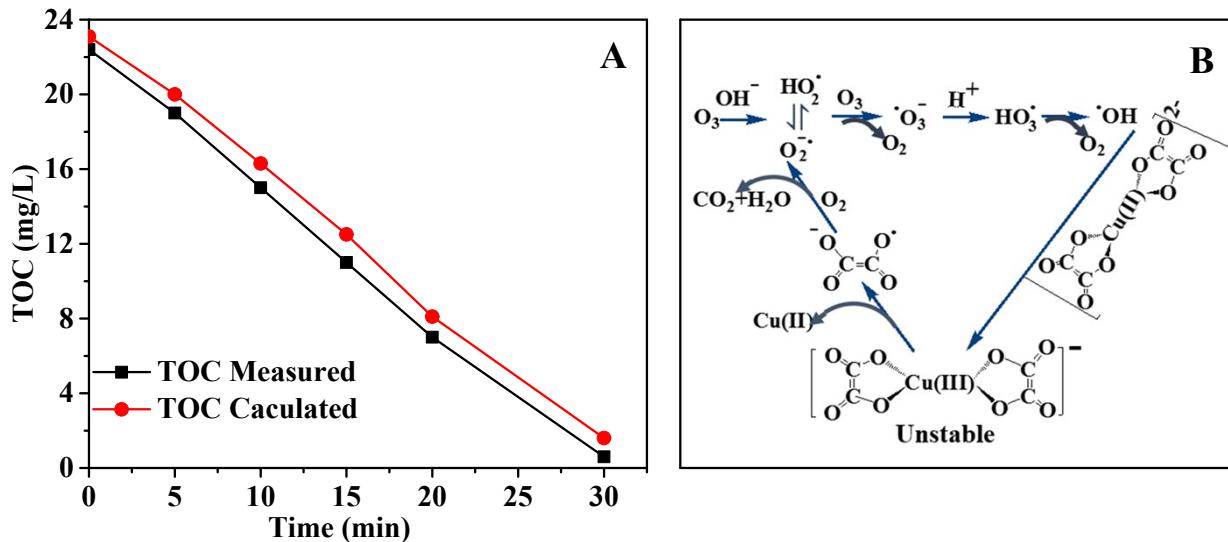
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134 Figure S7. (A) Measured TOC vs. Calculated TOC based on OA concentration. [Initial OA: 1.0
 135 mM; working solution: 200 mL; catalyst dose: 4.5 mg/L Cu^{2+} ; pH = 6.59 ± 0.20]. (B) Proposed
 136 reaction mechanism of OA degradation during homogeneous catalytic ozonation with copper ion
 137 [Note: (1).Only representative Cu-oxalate complex is shown; and (2). Results shown in Figure
 138 S7A indicate the intermediate products are either unstable (e.g. radicals) or indistinguishable
 139 from oxalate (e.g. Cu-oxalate complex) using the current analytical method (HPLC-DAD)].

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